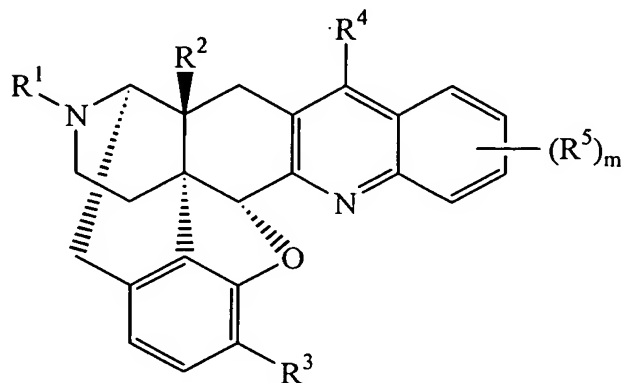


CLAIMS

1. A therapeutic or prophylactic agent for frequent urination or urinary incontinence, comprising as an effective ingredient a quinolinomorphinan derivative of the Formula (I):



(I)

5 (wherein R^1 is hydrogen, C_1 - C_5 alkyl, C_4 - C_7 cycloalkylalkyl, C_5 - C_7 cycloalkenylalkyl, C_6 - C_{12} aryl, C_7 - C_{13} aralkyl, C_2 - C_7 alkenyl, C_1 - C_5 alkanoyl, furan-2-ylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5), or thiophene-2-ylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5);

10 R^2 and R^3 independently are hydrogen, hydroxy, C_1 - C_5 alkoxy, C_1 - C_5 alkanoyloxy, C_7 - C_{13} aralkyloxy or C_7 - C_{13} arylcarbonyloxy;
 m is an integer of 0 to 4;

$R^5(s)$ is(are) substituent(s) on the benzene ring and independently represent(s) R^{18} , or two R^5 s on adjacent carbon atoms cooperatively represent a fused ring structure A (with the proviso that the remaining 0 to 2 R^5 s independently represent R^{18} or a pair of R^5 s represent another fused ring structure A),

wherein said ring structure A is benzo, indeno, naphtho or pyrido, or C_5 - C_7 cycloalkeno, each of which is substituted with 0 to 4 R^9 s, or non-substituted dioxoleno;

20 R^9 and R^{18} represent (1) independently fluoro, chloro, bromo, iodo, nitro, hydroxy,

C₁-C₅ alkyl, C₁-C₅ alkoxy, isothiocyanato, trifluoromethyl, trifluoromethoxy, cyano, phenyl, C₁-C₃ hydroxyalkyl, SR⁶, SOR⁶, SO₂R⁶, (CH₂)_kCO₂R⁷, SO₂NR⁷R⁸, CONR⁷R⁸, (CH₂)_kNR⁷R⁸, or (CH₂)_kN(R⁷)COR⁸ (wherein k is an integer of 0 to 5, R⁶ is C₁-C₅ alkyl, R⁷ and R⁸ independently are hydrogen, C₁-C₅ alkyl or C₄-C₆ cycloalkylalkyl), and/or (2) R⁹ and R¹⁸ on adjacent carbon atoms via the fused ring moiety cooperatively form R⁹-R¹⁸ which represents a bridging structure selected from the group consisting of ethano, propano and *o*-benzeno;

R⁴ is hydrogen, C₁-C₅ alkyl, C₁-C₅ hydroxyalkyl, C₆-C₁₂ aryl (which may be substituted with one or more substituents R¹⁷), NR¹⁰R¹¹, OR¹², COOR¹³ or CONR¹⁴R¹⁵, or R⁴ and R⁵ which substitutes on the peri-position cooperatively form R⁴-R⁵ that represent a bridging structure selected from the group consisting of N(R¹⁶)CO, N(R¹⁶)C(=NH), N(R¹⁶)CH₂, *o*-benzeno, ethano, propano and butano; R¹⁷ is fluoro, chloro, bromo, iodo, nitro, amino, hydroxy, C₁-C₅ alkyl, C₁-C₅ alkoxy, C₁-C₅ alkanoyloxy, trifluoromethyl, trifluoromethoxy or cyano; R¹⁰, R¹¹, R¹² and R¹⁶ independently are hydrogen, C₁-C₅ alkyl, C₄-C₇ cycloalkylalkyl, C₇-C₁₃ aralkyl or C₁-C₅ alkanoyl, and R¹³, R¹⁴ and R¹⁵ independently are hydrogen, C₁-C₅ alkyl, C₆-C₁₂ aryl or C₇-C₁₃ aralkyl)

or a pharmaceutically acceptable acid addition salt thereof.

2. A therapeutic or prophylactic agent for frequent urination or urinary incontinence according to claim 1, wherein in Formula (I), R⁴ is hydrogen, C₁-C₅ alkyl or NR¹⁰R¹¹, or R⁴ and R⁵ which substitutes on the peri-position cooperatively form R⁴-R⁵ that represent a bridging structure N(R¹⁶)CO (wherein R¹⁰, R¹¹ and R¹⁶ represent the same meanings as in claim 1).

3. A therapeutic or prophylactic agent for frequent urination or urinary incontinence according to claim 2, wherein in Formula (I), R¹ is C₁-C₅ alkyl, C₄-C₇ cycloalkylalkyl, C₇-C₁₃ aralkyl, furan-2-ylalkyl (wherein the number of carbon atoms in the alkyl moiety is 1 to 5), or thiophene-2-ylalkyl (wherein the number of carbon

atoms in the alkyl moiety is 1 to 5).